● PRINTER RUSH ● (PTO ASSISTANCE)

Application :	09/76/30/37	Examiner :	Low	GAU:	1614 ulglad
From:	the	Location: (DC) FMF FDC	Date:	1118104
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[RUSH] MES	sage: In to ta. Campt While.	1.1.	ce "c" of Ta twing bolded	ble 1 it a and non-be	cless to clad data.
[XRUSH] RE	sponse:	de pointed terrieu of is bolded	cut ky a n Feb 2,	aplicant 2005, M	- during
In a	ddition this	rousd appears.	to be an o should be	msue fr resc <i>ann</i> INITI	PROBLEM ALS: AM

NOTE: This form will be included as part of the official USPTO record, with the Response document coded as XRUSH.

REV 10/04

		TABL	Е1. A	TABLE 1. AZT/3TC prophylaxis of cats starting 3 days before FIV inoculation) prophy	laxis of	cats sta	rting 3	days bef	ore FIV	' inocula	tion				
	AZT/3TC treatment ^a				FIV levelsh	ls ^h										
	(kg/mg)	FIV		>	VI/PCR/vRNA	RNA			FI	FIV antibodies ^b	Jies ^b			CD4/C	CD4/CD8 ratio**	8 1
Cat #	-0.4-4-5-6-7 wk	inocul. Pre	Pre	4 wk	9 wk	II wk	4 wk 9 wk 11 wk 14 wk Pre	Pre	4 wk	9 wk	4 wk 9 wk 11 wk 14 wk	14 wk	Pre	7 wk	7 wk 11 wk 14 wk	14 wk
DHS	75-34-10	+	+		++-	++-	-1-1-			'			3.30	2.86	2.62	2.38
3GB	75-34-10-0-5	+	-/-/-	-/-/-	-/-/-	-1-1-	-/-/-	•	•	•	•	•	1.56	1.37	1.37	1.47
RU2	75-34-10	+	-/-/-	-/-/-	-/-/-	-/-/-	<i>+</i>	•	•	•	•	•	1.77	1.21.	1.18	- T
RUI	75-34-10	•	-/-/-	-/-/-	-1-1-	-/-/-	-1-1-	•	•		1	•	2.37	1.62	1.62	1.47
NK4	•	+	-/-/-	+/+/+	+/+/+	+/+/+	+/+/+	•	•	+	+	+	1.82	1.55	0.96	(16:0) (16:0)
NK6	•	+	-/-/-	+/+/-	+/+/+	+/+/+	+/+/+	•	1	+	. +	+	1.61	0.92	1	(C.4.5)
101	•	+	-/-/-	-/+/+	-/+/+	-/+/+	-/+/+	•	٠,	+	+	+	3.40) 5. j	<u> </u>
IH5	•	+	-/-/-	+/+/-		+/+/+ +/+/+	+/+/+	٠	,	+	+	+	4.40	1.34	09:0	(1.6 <u>1</u>

⁴ The AZT/3TC treatment was started 3 days before FIV inoculation (-0.4 post-infection) at a dosc of 75 mg/kg each and decreased to 34 mg/kg at 4 wk postinfection (pi) and then to 10 mg/kg at 5 wk pi. In one cat (#3GB), the AZT/3TC treatment was withdrawn at 6 wk pi and resumed at a low dose of 5 mg/kg at 7 wk pi. The changes in doses if each drug, including the amount (mg/kg) and time (wk pi), are shown.

by immunoblot analysis. In general, RT-PCR for plasma viral RNA was less sensitive than PCR of FIV provirus in PBMC after amplification of infected cells ^b Samples before drug or placebo treatment (Pre) and those at various weeks post-infection (wk) were tested for FIV levels, FIV antibodies, and CD4/CD8 rations. FIV levels were determined by virus isolation (VI), PCR for FIV provirus in PBMC, and RT-PCR for plasma viral RNA (vRNA). FIV untibodies were determined by coculturing.

Inverted CD4/CD8 ratios are bolded